AMENDMENTS TO THE CLAIMS

The following is a complete listing of the claims, which replaces all previous versions and listings of the claims.

1-35. (Cancelled)

36. (Previously Presented) Chelating agent of the general formula:

wherein m is 0 or 1;

X is NR4 or S;

Y is SR₅, NHR₅ or P(R₅)₂;

R₁ and R₃ are the same or different and are selected from H, alkyl or aryl;

R2 is H, COOH, NHR6 or (CH2)nCOOR6;

R₄ is H, alkyl, aryl, (CH₂)_nCOOR₆ or (CH₂)_nOR₆;

R₅ is H, alkyl, aryl, (CH₂)_nCOOR₆ or (CH₂)_nOR₆;

R₆ is H, alkyl or aryl;

n is 1, 2, 3, 4, 5, 6, 7, 8, 9 or 10; and

wherein at least one of R_1 , R_3 , R_4 , R_5 , and R_6 is phenyl or benzyl.

37. (Cancelled)

38. (Currently Amended) Chelating agent of the general formula:

wherein m is 0 or 1;

X is NR4 or S;

Y is SR₅, NHR₄ or P(R₅)₂;

 R_1 and R_3 are the same or different and are selected from H, alkyl or aryl, wherein at least one of R_1 and R_3 is aryl;

R2 is H, COOH, NHR6 or (CH2)nCOOR8;

R₄ is H. alkvl. arvl. (CH₂)_nCOOR₆ or (CH₂)_nOR₆;

Rs is H, alkyl, aryl, (CH2)nCOOR6 or (CH2)nOR6;

R₈ is H, a biomolecule, alkyl or aryl; and

n is 1, 2, 3, 4, 5, 6, 7, 8, 9 or 10.

39-42. (Cancelled)

- 43. (Previously Presented) A metal complex comprising the chelating agent of claim 36.
- 44. (Previously Presented) Chelating agent as claimed in claim 38, wherein R_{θ} is a biomolecule.
- 45. (Previously Presented) Chelating agent as claimed in claim 44, wherein the biomolecule is selected from amino acids, peptides, proteins, oligonucleotides, polynucleotides, and sugars.
- 46. (Previously Presented) Chelating agent as claimed in claim 44, wherein the biomolecule is selected from the group consisting of antibodies and ligands of tumor receptors.

- 47. (Previously Presented) Chelating agent as claimed in claim 44, wherein the biomolecule is selected from the group consisting of CCK, thioglucose, glucosamine, somatostatin, neurotensin, bombesin, annexin, interleukins, growth factors, steroid hormones and molecules binding to GPIIb/IIIIa receptors.
- 48. (Previously Presented) Chelating agent as claimed in claim 44, wherein the biomolecule is selected from the group consisting of glucose, thioglucose, and neurotransmitters.
- 49. (Previously Presented) Chelating agent as claimed in claim 44, wherein the biomolecule is an inhibitor of the tyrosine kinase activity.
 - 50. (Cancelled)
- 51. (Previously Presented) The chelating agent as claimed in claim 36, wherein when R_1 = R_3 = CH_3 , R_2 , R_4 and R_6 are not all three H.
- 52. (Previously Presented) The chelating agent as claimed in claim 38, wherein when R_1 or R_3 is CH₃, R_2 , R_4 and R_5 are not all three H.
- 53. (New) Chelating agent as claimed in claim 36, wherein alkyl is a C_1 alkyl, C_2 alkyl, C_3 alkyl, C_4 alkyl, C_5 alkyl, C_6 alkyl, C_8 alkyl, C_8 alkyl, C_9 alkyl,
- 54. (New) Chelating agent as claimed in claim 53, wherein alkyl is methyl, ethyl, n-propyl, isopropyl, n-butyl, isobutyl, s-butyl, h-butyl, n-pentyl, isopentyl, neopentyl, n-hexyl, isohexyl (2-methylpentyl), neohexyl (2,2-dimethylbutyl), 3-methylpentyl, 2,3-dimethylbutyl.
 - 55. (New) Chelating agent as claimed in claim 36, wherein n is 2, 3, 4, 5 or 6.
- 56. (New) Chelating agent as claimed in claim 38, wherein alkyl is a C_1 alkyl, C_2 alkyl, C_3 alkyl, C_4 alkyl, C_5 alkyl or C_6 alkyl.
- 57. (New) Chelating agent as claimed in claim 56, wherein alkyl is methyl, ethyl, n-propyl, isopropyl, n-butyl, isobutyl, s-butyl, t-butyl, n-pentyl, isopentyl, neopentyl, n-hexyl, isohexyl (2-methylpentyl), neohexyl (2,2-dimethylbutyl), 3-methylpentyl, 2,3-dimethylbutyl.
 - 58. (New) Chelating agent as claimed in claim 38, wherein n is 2, 3, 4, 5 or 6.